Note 3: The subject of this AD is addressed in Direction Generale De L'Aviation Civile (France) AD 94–077–016(B)R1 and AD 94–076–036(B)R1, both dated December 4, 1996.

Issued in Fort Worth, Texas, on February 26, 1998.

Eric Bries,

Acting Manager, Rotorcraft Directorate, Aircraft Certification Service.

[FR Doc. 98–5733 Filed 3–5–98; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 98–ANE–92]

Amendment to Class E Airspace; Laconia, NH; Correction

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Direct final rule; correction.

SUMMARY: This action corrects a charting error in the description of revised Class E airspace at Laconia, NH (KLCI) published in the **Federal Register** on February 20, 1998 (63 FR 8563) and intended to provide adequate controlled airspace for those aircraft using the new GPS RWY 26 standard instrument approach procedure to Laconia Municipal Airport.

DATES: Effective 0901 UTC, April 23, 1998.

Comments for inclusion in the Rules Docket must be received on or before March 23, 1998.

ADDRESSES: Send comments on the rule to: Manager, Airspace Branch ANE–520, Federal Aviation Administration, Docket No. 98–ANE–92, 12 New England Executive Park, Burlington, MA 01803–5299; telephone (781) 238–7520; fax (781) 238–7596. Comments may also be sent electronically via the internet to the following address: ''9 ne airspacefaa.dot.gov''. Comments sent electronically must indicate Docket 98–ANE–92 in the subject line.

The official docket file may be examined in the Office of the Regional Counsel, New England Region, ANE–7, Room 401, 12 New England Executive Park, Burlington, MA 01803–5299; telephone (781) 238–7050; fax (781) 238–7055.

An informal docket may also be examined during normal business hours in the Air Traffic Division, Room 408, by contacting the Acting Manager, Airspace Branch at the first address listed above.

FOR FURTHER INFORMATION CONTACT:

David T. Bayley, ANE–520.3, 12 New England Executive Park, Burlington, MA 01803–5299; telephone (781) 238–7523; fax (781) 238–7596.

SUPPLEMENTARY INFORMATION: On February 20, 1998, the FAA published in the **Federal Register** a direct final rule revising the Class E airspace at Laconia, NH (KLCI) to provide for adequate controlled airspace for those aircraft using the new GPS RWY 26 standard instrument approach procedure to Laconia Municipal Airport (63 FR 8563). Since publication of that direct final rule, the FAA has been advised of a charting error in the description of the Class E airspace at Laconia. This action corrects that error.

Correction to the Direct Final Rule

Accordingly, pursuant to the authority delegated to me, the amendment to Class E airspace at Laconia, NH as published in the **Federal Register** on February 20, 1998 (63 FR 8563), **Federal Register** document 98–4314; and the description in FAA Order 7400.9E, dated September 10, 1997, and effective September 16, 1997, which is incorporated by reference in 14 CFR 71.1 are corrected as follows:

§ 71.1 [Corrected]

On page 8564, column 3, 9th and 10th lines, correct the words ''Belknap NDP 249° bearing'' to read ''Belknap NDB 249°/069° bearings''.

Issued in Burlington, MA, on February 26, 1998.

Bill Peacock,

Manager, Air Traffic Division, New England Region.

[FR Doc. 98–5693 Filed 3–5–98; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 173

[Docket No. 97F–0038]

Secondary Direct Food Additives Permitted in Food for Human Consumption

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of acidified solutions of sodium chlorite as an antimicrobial agent in the processing of red meat. This action is in response to a petition filed by Alcide Corp.

DATES: This regulation is effective March 6, 1998; written objections and requests for a hearing by April 6, 1998. The Director of the Office of the Federal Register approves the incorporation by reference in accordance with 5 U.S.C. 552(a) and 1 CFR part 51 of certain publications in § 173.325(d) (21 CFR 173.325(d)), effective March 6, 1998.

ADDRESSES: Written objections may be sent to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Robert L. Martin, Center for Food Safety and Applied Nutrition (HFS–217), Food and Drug Administration, 200 C St. SW., Washington, DC 20204–0001, 202–418–3074.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of February 5, 1997 (62 FR 5428), FDA announced that a food additive petition (FAP 7A4532) had been filed by Alcide Corp., Inc., 8561 154th Ave. NE., Redmond, WA 98052, proposing that the food additive regulations be amended to provide for the safe use of acidified sodium chlorite solutions for red meat disinfection in processing plants. In its evaluation of the petition, the agency has concluded that red meat is not disinfected, but that the microbial contamination of the meat is reduced. Therefore, the agency is approving this additive as an antimicrobial agent in red meat processing.

FDA has evaluated data in the petition and other relevant material. The agency has also consulted with scientists from the Food Safety and Inspection Service, U. S. Department of Agriculture, concerning the technological and practical aspects of the proposed use of acidified sodium chlorite solutions. Based upon this information and consultation, the agency concludes that the proposed use of the additive is safe, and the additive will have the intended technical effect of reducing microbial contamination on red meat. Therefore, § 173.325 is being amended as set forth below. Additionally, the agency is revising § 173.325 to eliminate redundancy. This revision is strictly editorial and is not a substantive change in the regulation.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person